# VOC Capture and Destruction Efficiency Test Protocol

Ferrara Candy Company
Four Permanent Total Enclosures and
Catalytic Oxidizer System
Forest Park, Illinois
Protocol No. M161403



# **VOC Destruction Efficiency Test Protocol**

# Ferrara Candy Company Four Existing Permanent Total Enclosures and Catalytic Oxidizer System Forest Park, Illinois

Protocol No. M161403

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Submitted By

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Sample Point Location Diagrams
Test Train Diagrams
Data Sheets
Calculation and Nomenclature Sheets
Calibration Data

# 1.0 INTRODUCTION

An emission compliance test program to determine the volatile organic compounds (VOC) capture and destruction efficiency and emission rate of the four existing Permanent Total Enclosures (PTEs) and the Catalytic Oxidizer System which controls Candy polishing process emissions will be conducted at the Ferrara Pan Candy Company in Forest Park, Illinois. All testing will be performed as described in the *Code of Federal Regulations*, Title 40, Part 60, Appendix A (40CFR60), Methods 1, 2, 3, 4, 18 (methane only, if applicable), 25A, and 40CFR, Part 51, Appendix M, Method 204, and ASTM E337-02 and the latest revisions thereof. Where applicable, the *Quality Assurance Handbook for Air Pollution Measurement Systems*, Volume III, Stationary Source Specific Methods, USEPA 600/4-77-027b will be used to determine the precise procedures.

Project Contact Information						
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# 2.0 PROCESS DESCRIPTION AND OPERATING RATES

#### **Process Description and Rates**

The candy polishing process is a batch process in which candy centers are placed into ovate rotating pans where the pans are filled with a pre-weighed amount of centers. The typical batch process from center addition to pan unloading takes approximately one hour. After the centers are added, the chocolate or other "grossing" raw materials are added by hand and until the desired size of the candy pieces is obtained. Once the candy is sized properly, a pre-measured amount of confectioner's glaze is added by hand. Once added, the pans rotate until the glaze is dried and the batch is then finished and ready for packaging.

The glazes are either vegetable/mineral oil or solvent-based. The solvent-based glazes consist mainly of solids dissolved in ethanol, ethyl acetate, or other minor food grade solvents that evaporate to leave a hardy, shiny coating on the candy. Vegetable/mineral oil-based glazes act in a similar fashion but dry more slowly.

Candies polished include chocolate, jelly beans, and other soft candies as described below:

<u>Chocolate Panning/Polishing</u> usually starts with almond, peanut, or raisin centers rather than sugar centers as is used for the jelly beans and soft candy panning process. Steam heated chocolate is sprayed or poured onto the centers as the pans rotate and conditioned air solidifies the chocolate as the centers are covered and increased in size. After the correct amount of chocolate is applied, a confectioner's glaze, which may contain VOCs, is hand poured in premeasured amounts once per batch onto the candy pieces. The solvent portion of the glaze evaporates and leaves a hard, glossy shine (e.g., polish) to the candy pieces. The chocolate

panning/polishing process in conducted in permanent total enclosures (PTEs) and VOC emissions are collected and routed to a catalytic oxidizer for destruction.

Once the candy glaze solidifies, the batch is complete and the candy pieces are removed from the pans, placed onto trays and transferred to packaging.

<u>Soft Candy Panning/Polishing</u> usually starts with "grossed" candy centers and other ingredients such as sugar, colors, and flavors which are added to build up a shell around the centers. As a final step, a confectioner's glaze, which may or may not contain VOCs, is added once per batch by hand in pre-measured amounts as a protective coating and decorative layer. The soft candy panning/polishing process in conducted in a PTE and VOC emissions are collected and routed to a catalytic oxidizer for destruction.

As with the chocolate polishing process, once the glaze solidifies the batch is completed and the candy pieces are removed from the pans, placed onto trays and transferred to packaging.

#### **Stack Testing Operating Parameters**

During the stack testing program, all polishing pans will be in operation in the PTEs. The testing will be conducted during the entire batching process from center addition through glaze application and drying. Each pan will polish a typical batch size of approximately 250 lbs of candy pieces and use the confectioner's glaze in the prescribed pre-measured quantity for each pan which is typically about 8 ounces of glaze.

# 3.0 TEST REQUIREMENTS

The following table presents a list of the pollutants to be tested at each emission source and the applicable rules or regulations for each emission test:

TEST PARAMETERS						
<b>Emission Point</b>	Pollutant Tested	Method/Regulation Citation				
Catalytic Oxidizer Inlet	VOC	USEPA Methods 1, 2, 3, 4, 18 (methane only, if needed), 25A, 40CFR60, Appendix A and ASTM E337-02				
Catalytic Oxidizer Stack	VOC	USEPA Methods 1, 2, 3, 4, 18 (methane only, if needed), 25A, 40CFR60, Appendix A and ASTM E337-02				
Permanent Total Enclosures	VOC	USEPA Method 204, 40CFR51, Appendix A.				

# **VOC Destruction Efficiency Tests**

Three (3) one (1)-hour VOC test runs will be performed simultaneously at the catalytic oxidizer inlet and stack in accordance with United States Environmental Protection Agency (USEPA) Methods 1, 2, 3, 4, 18 (methane only, if applicable), and 25A, Title 40, *Code of Federal Regulations*, Part 60, Appendix A, and ASTM Method E337-02 wet bulb/dry bulb procedures.

If required, one integrated Tedlar bag sample will be taken during each test run and analyzed off-site for methane concentrations by gas chromatograph/flame ionization detector for total gaseous non-methane hydrocarbon determinations.

Destruction Efficiency (DE) will be calculated as follows:

$$DE = \frac{Oxidizer InVOClb/hr - Oxidizer OutVOClb/hr}{Oxidizer InVOClb/hr} \times 100$$

Oxidizer Test Locations							
Location	Dimensions	Area	Test Parameter	Number of Sampling Points			
Oxidizer Inlet	30 Inches	4.909 Square Feet	Volumetric flow	16			
Oxidizer Outlet	31 inches	5.241 Square Feet	Volumetric flow	16			

# 4.0 SPECIFIC TEST PROCEDURES

Detailed test procedures are appended. Three complete test runs will be performed for each parameter in accordance with the following USEPA methods.

- Volumetric flow will be determined utilizing USEPA Methods 1 and 2, 40CFR60 in conjunction with each emission test. A pretest and posttest volumetric flow will be performed and averaged for each VOC gaseous test run.
- 2. Oxygen (O<sub>2</sub>) and carbon dioxide (CO<sub>2</sub>) content of the gases will be determined by USEPA Method 3, 40CFR60 during each test run.
- 3. Moisture content of the gases will be determined using USEPA Method 4, 40CFR60 and/or ASTM E337-02 during each volumetric flow test run.
- 4. Total hydrocarbon emissions will be determined in accordance with USEPA Method 25A, 40CFR60. Each test will be one hour long with the emissions determined as ppm propane.
- 5. If required, one integrated Tedlar bag sample will be taken during each hour of testing at each test location during each test run and analyzed off-site for methane concentrations by gas chromatograph/flame ionization detector for total gaseous non-methane hydrocarbon determinations.
- 6. The four Permanent Total Enclosures (PTE's) will be evaluated to verify that they meet the criteria of USEPA Method 204, 40CFR51, Appendix M.

7. Where applicable, the *Quality Assurance Handbook for Air Pollution Measurement Systems*, Volume III, Stationary Source Specific Methods, USEPA 600/4-77-027b is used to determine the precise procedures.

# 5.0 TEST PROGRAM SCHEDULE

Refer to plant submittals for specific dates.

Day	Activity	On-Site Hours
1	Set-up equipment and evaluate four PTEs.	8
2	Perform VOC destruction efficiency test runs.	8

All project days are considered weekdays and are scheduled between the hours of 7:00 a.m. and 6:00 p.m., unless otherwise stated.

# **6.0 PROJECT PERSONNEL**

- 1 Project Manager
- 2 Test Engineers
- 1 Test Technician

# 7.0 PLANT REQUIREMENTS

Mostardi Platt must be supplied with the following items in order to complete this test program:

- 1. Safe access to test positions.
- 2. Electrical power 110 V, 20 A service at the test locations.
- 3. Test ports cleaned and loose prior to arrival of test crew.
- 4. Any scaffolding required to perform the tests.
- 5. Sufficient lighting at the test site.
- 6. Plant or pollution control equipment-operating data, if required for report.
- 7. Washroom facilities for use by members of the test crew.
- 8. Steady load during test period.
- 9. Communication between the test location and the control room.
- 10. Parking location to place Mostardi Platt mobile trailer within 200 feet of sampling locations with access to multiple 110 V, 15 amp, 60-cycle, 220 V, 50 amp, 60-cycle circuits, or a 480 V welding receptacle.
- 11. Oxidizer operating temperatures should be recorded during testing and provided to test crew for final report.

# 8.0 TEST PROCEDURES

All testing, sampling, analytical, and calibration procedures used for this test program will be performed as described in the Title 40, *Code of Federal Regulations*, Part 60, Appendix A (40CFR60), Methods 1, 2, 3, 4, 25A, and, 40CFR51, Appendix M, Method 204, and ASTM E337-02 and the latest revisions thereof. Where applicable, the *Quality Assurance Handbook for Air Pollution Measurement Systems*, Volume III, Stationary Source Specific Methods, United States Environmental Protection Agency (USEPA) 600/R-94/038c, September 1994 was used to supplement procedures.

## **Volumetric Flowrate Determination**

In order to determine the VOC emission rates on a lb/hr basis, the gas velocities and volumetric flowrates are determined using reference Method 2.

Velocity pressures are determined by traversing the test locations with S-type pitot tubes. Temperatures are measured using a K-type thermocouple with a calibrated digital temperature indicator. The molecular weight and moisture content of the gases are determined to permit the calculation of the volumetric flowrate. Sampling points utilized are determined using Method 1, 40CFR60.

# Oxygen (O<sub>2</sub>)/Carbon Dioxide (CO<sub>2</sub>) Determination

 $O_2$  and  $CO_2$  gas content are determined at the test locations in accordance with Method 3, 40CFR60. This method collects samples in a grab or integrated manner and analyzes the samples using a Fyrite gas analyzer. Several passes of the gas are made during each run to ensure a stable reading. Mandatory leak checks are performed prior to and following each use. Chemicals are changed frequently and inspected for reactivity prior to each use.

# Moisture (H<sub>2</sub>O) Determination

Determining the moisture content in the gas stream is necessary to calculate the stack gas volumetric air flow on a dry basis and the VOC emission rate in lb/hr. For this purpose, Mostardi Platt uses two methods.

1. American Standard Testing Method (ASTM) Method E337-02 reapproved 2002, wet bulb/dry bulb measurements are made at the oxidizer inlet and oxidizer outlet if temperatures allow during each VOC sampling run and the water vapor content is calculated as follows:

$$Bws = \left[\frac{e' - AP(t - t')}{P}\right]$$

where:

e' = saturated vapor pressure of water, in. Hg, at the wet bulb temperature, t'

 $A = 3.67 \times 10^{-4} [1 + 0.00064(t' - 32)]$ 

P = absolute pressure, in. Hg, in duct

t = dry bulb temperature, °Ft' = wet bulb temperature, °F

2. At the oxidizer outlet, if temperatures dictate, *An Alternative Method for Stack Gas Moisture Determination*, written by John Stanley and Peter Westlin, August 1978, Emission Measurement Branch, USEPA, is utilized. The sampling equipment is the same as specified for the moisture approximation method in Reference Method 4, except for the addition of two impingers, one containing silica gel.

Approximately 15 mls of water are added to each of the first two impingers and the third is left empty. An impinger containing approximately 15 grams of silica gel and a glass-wool-packed outlet is attached following the third impinger. The entire impinger train, excluding the inlet and outlet connectors, is weighed to the nearest 0.05 gram. The impingers are placed in an ice bath to maintain the sampled gas passed through the silica gel impinger outlet below 68°F in order to increase the accuracy of the sampled dry gas volume measurement. Each sample is extracted through a stainless steel probe at a constant sample rate of between one to four liters per minute, which is maintained during the course of the other simultaneous reference method sampling. After each run, a leak check of the sampling train is performed at a vacuum greater than the sampling vacuum to determine if any leakage had occurred during sampling. Following the leak check, the impingers are removed from the ice bath and allowed to warm. Any condensed moisture on the exterior is removed and the train reweighed.

# **Volatile Organic Compound (VOC) Determination**

The Method 25A sampling and measurement system meets the requirements for stack sampling of VOCs set forth by the United States Environmental Protection Agency (USEPA). In particular, it meets the requirements of USEPA Reference Method 25A, "Determination of Total Gaseous Organic Concentration Using a Flame Ionization Analyzer," 40CFR60, Appendix A. This method applies to the measurement of total gaseous organic concentration of hydrocarbons. With this method, gas samples are extracted from the sample locations through heated Teflon sample lines to the analyzers.

The flame ionization detectors (FIDs) to be used during this program are VIG High-Temperature Total Hydrocarbon Analyzers. They are highly sensitive FIDs that provide a direct reading of total organic vapor concentrations with linear ranges of 0-10, 100, 1000, and 10,000 ppm by volume. The instruments are calibrated using ultra-zero air and propane in air EPA Protocol standards. The calibrations are performed before and after sampling with calibration checks performed between each test run. Sampling is conducted continuously for three one-hour periods for VOC destruction efficiency tests. Sample times and locations are logged simultaneously on data loggers. Final concentrations are determined by subtracting methane from the Method 18 analysis, if applicable, from the inlet and outlet concentrations.

Calculations are performed by computer or by hand. An explanation of the nomenclature and calculations along with the complete test results is included in the appendix. Also appended are calibration data and copies of the raw field data sheets.

#### **Enclosure Evaluation Determination**

A 100% Permanent Total Enclosure (PTE) must meet four (4) specific engineering criteria. The criteria are described in USEPA Method 204, 40CFR51, Appendix M. A summary of these items and the evaluation technique used are described below.

#### Natural Draft Openings (NDO) Distance to Emitting Point

<u>Criteria</u>: All NDOs such as open doorways, windows, etc. must be at least four equivalent

NDO diameters from the nearest potential VOC emission point.

Technique: The dimensions of all NDOs and potential emission points are measured. The

calculated NDO equivalent diameters are compared to the emission point

distances measured.

#### **Total NDO Area**

Criteria: The area of all NDOs divided by the total area of all walls, floor and ceilings in the

enclosure (called the "NEAR" ratio in the procedure) must not exceed 0.05.

Technique: Actual measurements are used to determine a composite surface area of the

room and the NDOs and the NEAR ratio is determined.

#### **Velocity of Air Flow through NDOs**

Criteria: The calculated face velocity through the NDOs must be greater than 200 fpm.

This is defined as the total exhaust volume (in scfm), less make up air, divided by

the area of all NDOs (in square feet).

<u>Technique</u>: The composite area of the NDOs (as determined above) is used along with the

measured net exhaust volume, as determined by USEPA Method 2, 40CFR60, Appendix A, of the enclosure to calculate the face velocity or alternately, the static pressure of the enclosure is measured to determine if it meets the  $\geq 0.007$ 

inches H<sub>2</sub>O criteria.

#### **Direction of Air Flow through the NDO**

Criteria: The direction of air flow through all NDOs must be into the enclosure.

<u>Technique</u>: Smoke tubes are used at each NDO to measure the direction of the air flow. A

record of this data will be included on the scale drawing.

# 9.0 QUALITY ASSURANCE PROCEDURES

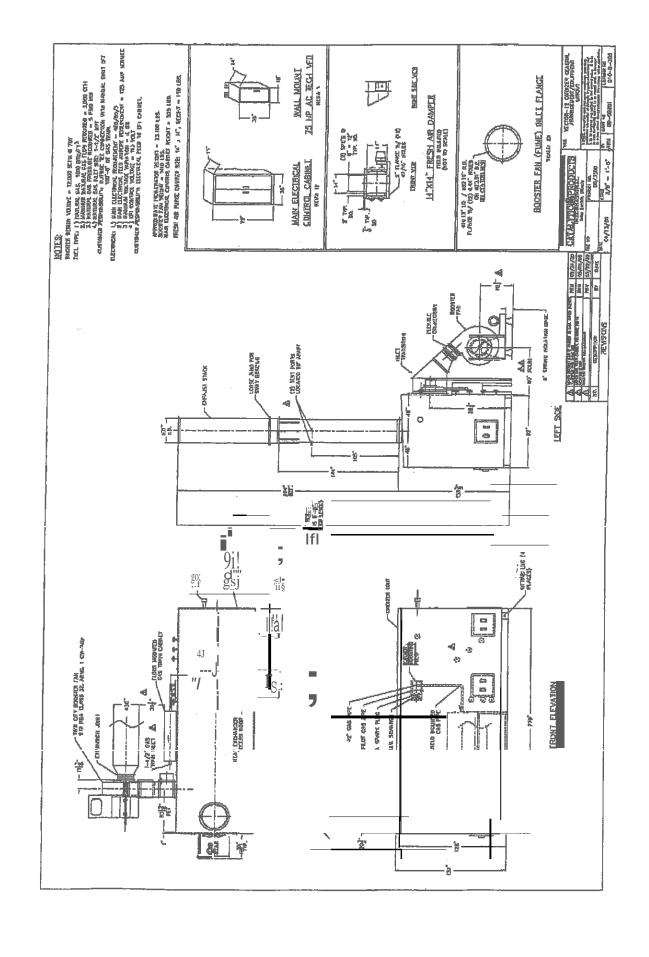
Mostardi Platt recognizes the previously described reference methods to be very techniqueoriented and attempts to minimize all factors which can increase error by implementing its Quality Assurance Program into every segment of its testing activities.

Shelf life of chemical reagents prepared at the Mostardi Platt laboratory or at the jobsite does not exceed those specified in the above mentioned methods. In addition, those reagents having a shelf life of one week are prepared daily at the jobsite. When on-site analyses are required, the same person performing the analysis performs all reagent standardization daily.

Dry and wet test meters are calibrated according to methods described in the *Quality Assurance Handbook for Air Pollution Measurement Systems*, Sections 3.3.2, 3.4.2 and 3.5.2. Percent error for the wet test meter according to the methods is less than the allowable error of 1.0%. The dry test meters measure the test sample volumes to within 2% at the flowrate and conditions encountered during sampling.

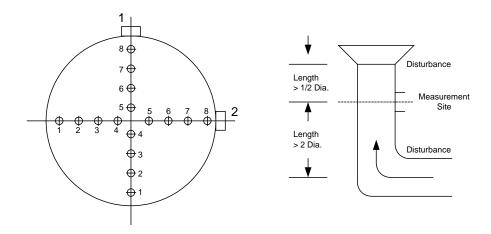
Calibration gases are EPA Protocol gases.

# **APPENDIX**





# **EQUAL AREA TRAVERSE FOR ROUND DUCTS**



Job: Ferrara Pan Candy Company

Forest Park, Illinois

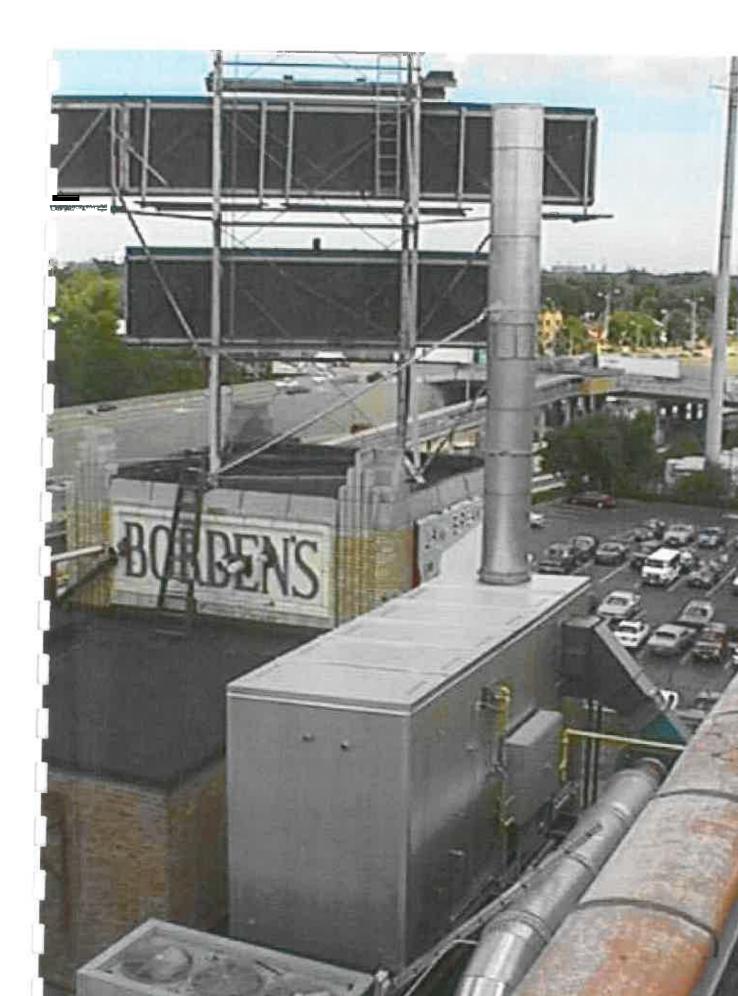
Test Location: Catalytic Oxidizer Inlet

Stack Diameter: 30 inches

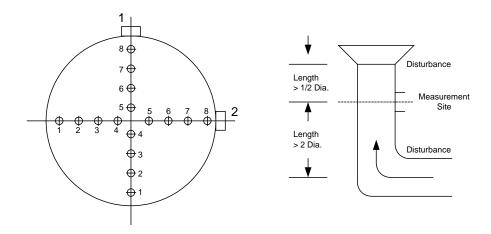
Stack Area: 4.909 Square Feet

No. Points Across Diameter: 8

No. of Ports: 2



# **EQUAL AREA TRAVERSE FOR ROUND DUCTS**



Job: Ferrara Pan Candy Company

Forest park, Illinois

Test Location: Catalytic Oxidizer Outlet

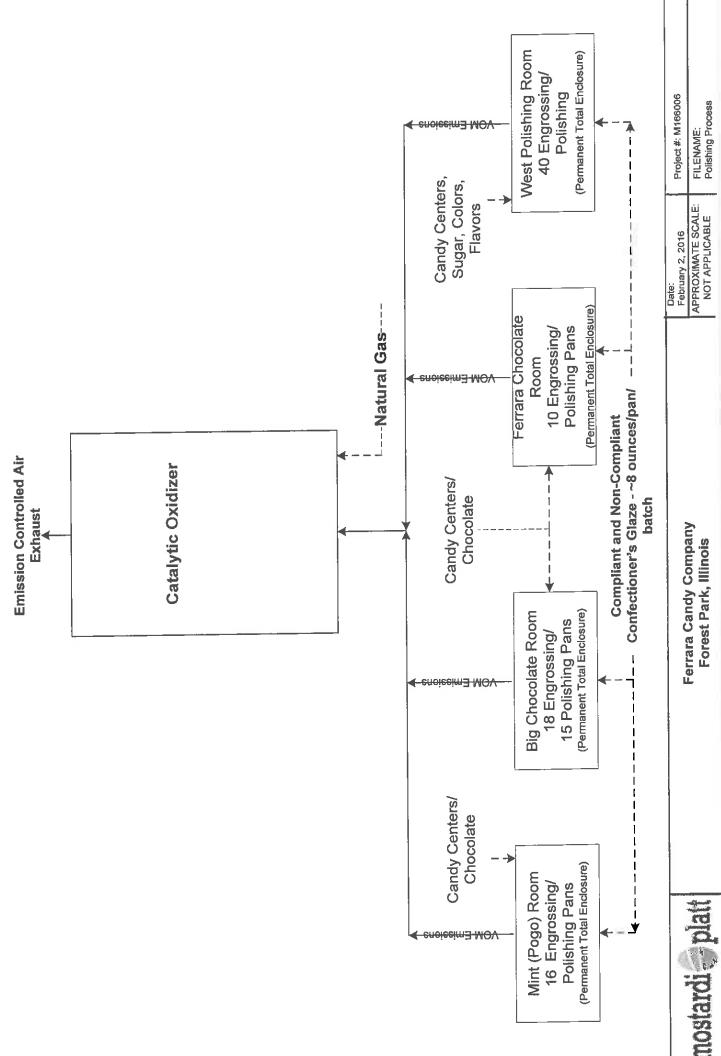
Stack Diameter: 31 Inches

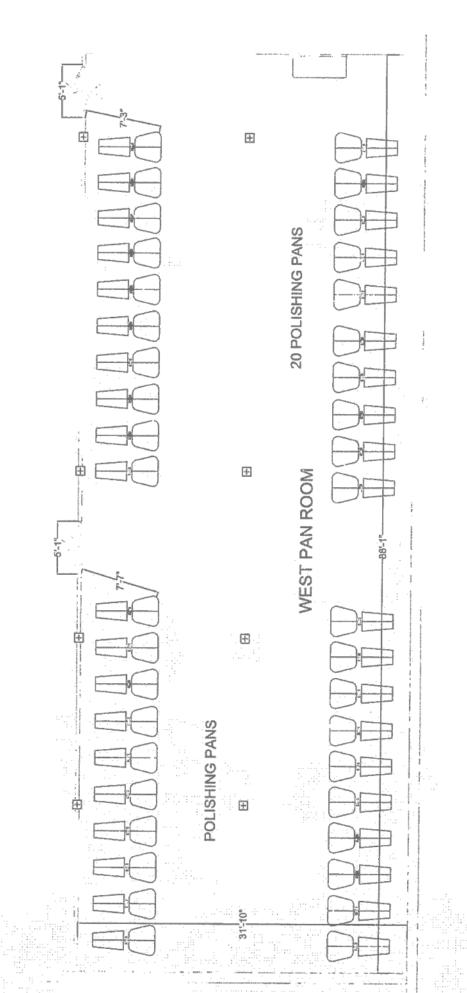
Stack Area: 5.241 Square Feet

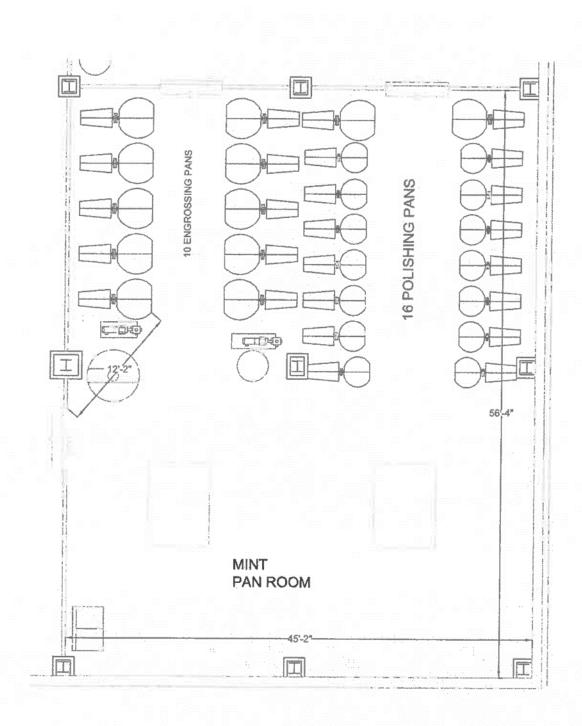
No. Points Across Diameter: 8

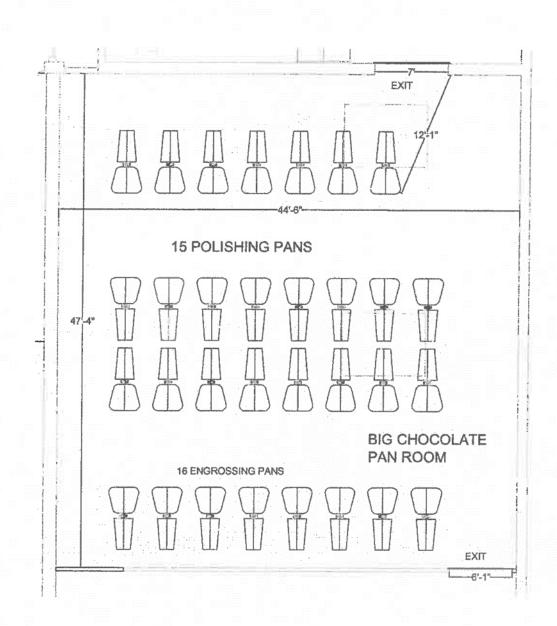
No. of Ports: 2

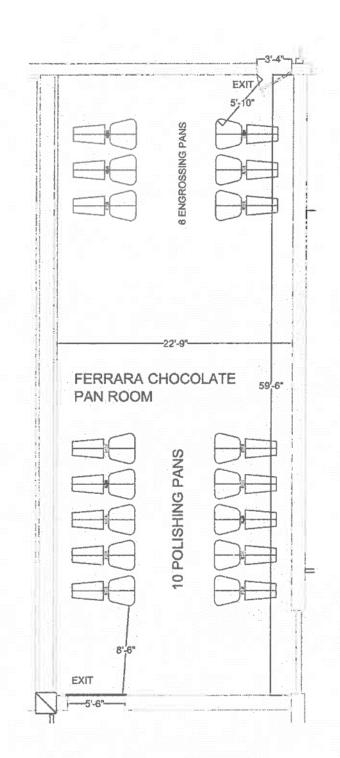
# Process Flow Diagram Catalytic Oxidation of Candy Polishing VOMs



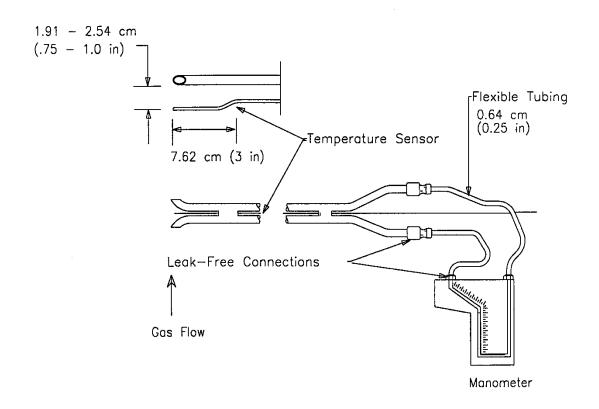




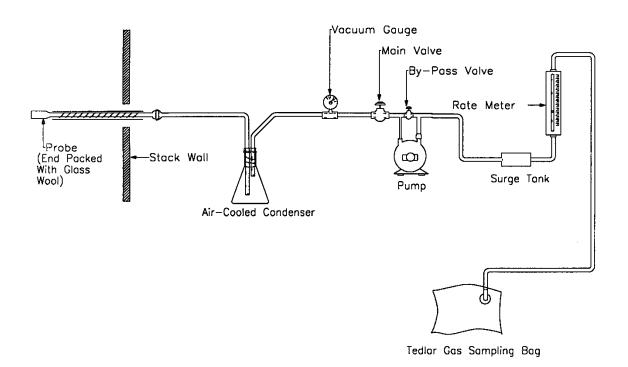




# **USEPA Method 2 - S-Type Pitot Tube Diagram**

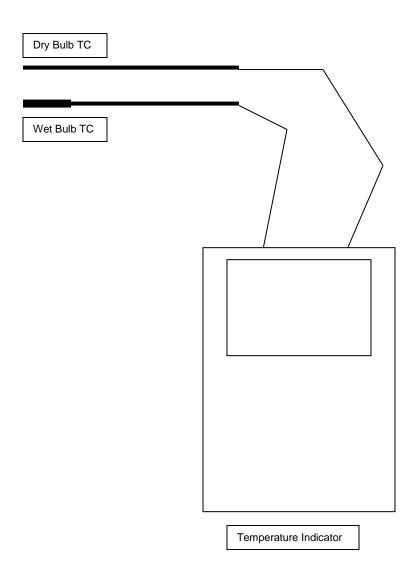


# **USEPA Method 3 - Integrated Oxygen/Carbon Dioxide Sample Train Diagram utilizing Fyrite Gas Analyzer**

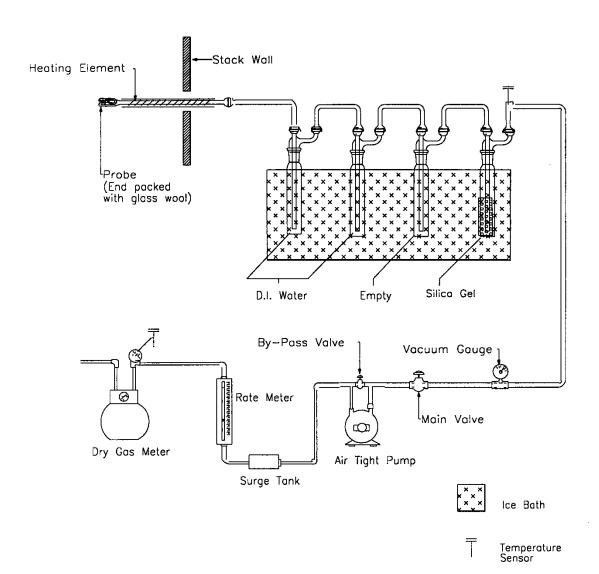


# **Moisture Determination**

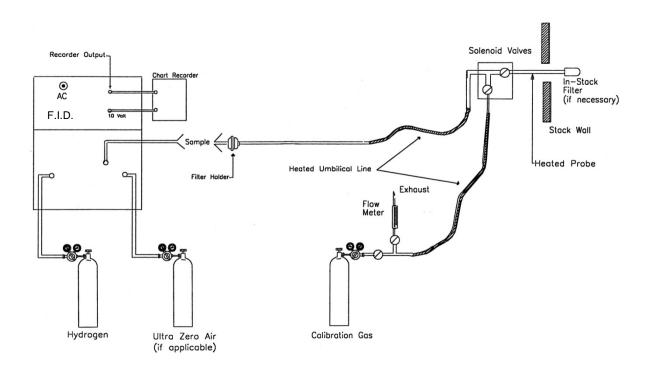
# **ASTM Method E337-02, Reapproved 2002**



# **USEPA Method 4 - Moisture Sample Train Diagram**



# **USEPA Method 25A -Total Gaseous Organic Compound Sample Train**



# **Volumetric Flow Rate Determination Field Data Sheet**

Project Nur	mber:				Dat	te:			
Client:					Tes	st Number	:		
Test Locati	on:				Sta	ırt Time:			
Source Cor	ndition:				End	d Time:			
Test Engine	eer:				Tes	st Tech:			
Flue Ar Port Le P <sub>bar</sub> Checks Static_ Static_	ea ngth ″Hg	CO2 %       Wet Bulb Temp        Leak         H2O O2 %       Dry Bulb Temp       Pre         Hg N2 %       B <sub>ws</sub> Post							
Port- Point #	ΔΡ	Temp. °F	$\sqrt{\Delta P}$	Null Point Angle, Degrees	Port- Point #	ΔР	Temp. °F	$\sqrt{\Delta P}$	Null Point Angle, Degrees
Average									
(	Md ×	$6 + .28 \times N_2\% = 1-Bws) + \sqrt{\frac{(}{}}$	(18 ×	Bws) =	_(Ms) √ <u>\ \ \ \ \ \ \ \ \ \ \ \ \ \ \ = \ \ \ \</u>	= ft/	/sec(Vs)		

\_\_\_\_\_ Vs × \_\_\_\_\_ Flue Area × 60 = \_\_\_\_\_acfm

 $17.647 \times \underline{\qquad} \operatorname{acfm} \times \frac{Ps}{Ts \, {}^{\circ}R} = \underline{\qquad} \operatorname{scfm} \times 60 = \underline{\qquad} \operatorname{scfh}$ 

# **MOISTURE FIELD DATA SHEET**

ProjectName/Number:		Date:
SamplingLocation:		
SourceCondition:		DryGasMeterNo.
Meter∆H	MeterY=	TestEngineer:

Test (Run)	No	Ba		essure (P <sub>bar</sub> )		_in. Hg	Orsat Analysis	
Gas Temperatu	ıre				in. Hg		%CO <sub>2</sub>	_%O <sub>2</sub>
Clock Time	Meter Volume (V <sub>m</sub> ) ft <sup>3</sup> or L	Meter Gage Pressure (ΔH)	Meter Temp. (t <sub>m</sub> )	Impgr. Outlet Temp °F	Condensate		Silica Gel or Train	
24 hour	(Circle One)	in. H <sub>2</sub> O	°F					
						mls (V <sub>f</sub> )	grams (W <sub>f</sub> ) grams (W <sub>i</sub> )	
	<u> </u>					mls	grams	
					$\times 0.04707 = \underline{\qquad}$ $ft^{3} [V_{wc(sto}]$	f	$\times 0.04715 = $	
					V <sub>m(std)</sub> =	ft <sup>3</sup>	= ft <sup>3</sup> [V <sub>w(std)</sub> ]	
					Water Vapor, prop Leak Check:	ortion by volun	ne B <sub>ws</sub> =	
							Moisture correction factor: 1 - B <sub>ws</sub> =	
Total Vol.	1				Comments:	L.	***	
Average				(T <sub>m</sub> )	°R			
Test (Run) Gas Temperatu	No	Ba°F Static Pre		essure (P <sub>bar</sub> )	in. Hg	in. Hg	Orsat Analysis %CO <sub>2</sub>	_%O <sub>2</sub>
Clock Time	Meter Volume (V <sub>m</sub> )	Meter Gage Pressure	Meter Temp.	Impgr. Outlet Temp	-	·		
24 hour	ft <sup>3</sup> or L (Circle One)	(ΔH) in. H <sub>2</sub> O	(t <sub>m</sub> ) °F	°F	<u>Condensate</u>		Silica Gel or Train	
	<del> </del>					1.00		
							grams (W <sub>f</sub> )	
					=	mls (V <sub>i</sub> ) mls	grams (W <sub>i</sub> ) grams	
						mls (V <sub>i</sub> ) mls fi	grams (W <sub>i</sub> )grams × 0.04715 =	
					 × 0.04707 =	mls (V <sub>i</sub> ) mls f <sub>i</sub> ] +f	grams (W <sub>i</sub> ) grams × 0.04715 =	
					ft^3 [V <sub>wc(stc)</sub> $V_{m(std)} =$	mls (V <sub>i</sub> ) mls fi] +f ft <sup>3</sup>	$ \begin{array}{cccccccccccccccccccccccccccccccccccc$	
					$\begin{array}{c} - \\ \times \ 0.04707 = \\ \underline{\qquad \qquad } ft^3 \ [V_{wc(std)} \\ \end{array}$	mls (V <sub>i</sub> ) mls fi] +f ft <sup>3</sup>	$ \begin{array}{cccccccccccccccccccccccccccccccccccc$	
					ft^3 [V <sub>wc(stc)</sub> $V_{m(std)} =$	mls (V <sub>i</sub> ) mls fi] +f ft <sup>3</sup>	grams (W <sub>i</sub> )grams × 0.04715 =  t <sup>3</sup> [V <sub>wsg(std)</sub> ] = ft <sup>3</sup> [V <sub>w(std)</sub> ]	
Total Vol.					ft^3 [V <sub>wc(stc)</sub> $V_{m(std)} =$	mls (V <sub>i</sub> ) mls fi] +f ft <sup>3</sup>	grams (W <sub>i</sub> )grams  × 0.04715 =  3 [V <sub>wsg(std)</sub> ] =ft <sup>3</sup> [V <sub>w(std)</sub> ]  ne  B <sub>ws</sub> =  Moisture correction factor:	

$$V_{m(std)} = 17.64 V_m Y \frac{P_{bar} + \frac{DH}{13.6}}{T_m}$$
 $B_{ws} = \frac{V_{w(std)}}{V_{w(std)} + V_{m(std)}}$ 

# **Method 25A Field Data Sheet**

Project:	Operator:		
Client:	Source:	1.	
Location:		2.	
Date:		3.	

Parameters	Location 1	Location 2	Location 3
Test 1			
Time			
VOC ppmv as			
Air flow, scfm			
VOC lbs/hr as			
Removal Efficiency, %			
Test 2			
Time			
VOC ppmv as			
Air flow, scfm			
VOC lbs/hr as			
Removal Efficiency, %			
Test 3			
Time			
VOC ppmv as			
Air flow, scfm			
VOC lbs/hr as			
Removal Efficiency, %			

# **METHOD 204 ENCLOSURE DATA SHEET**

Project: Sketch enclosure, all ducts, NDOs and potential						ential		
Location:				VOC emission points on accompanying page.				
Date:				Label all dimensions.				
Enclosure Designation:  Control Devices (s):		Process(e	Process(es) Enclosed:					
NDO to VC	C Emission Po	oint						
					Distances			

					Distances			
NDO	Dimensions	Area	Equivalent Diameter	VOC Emission Point	Minimum	Actual	Pass/ Fail?	

NDOs equivalent diameter = 
$$\left(\frac{4 \times \text{area}}{\pi}\right)^{0.5}$$

 $Minimum \ Allowed \ Distance = 4 \times Equivalent \ Diameter \ \ (NDO)$ 

NDO to Exhaust (TTE only)

						Distances		
Exhaust Point	Dimensions	Equivalent Diameter	NDO	Dimensions	Equivalent Diameter	Minimum	Actual	Pass/ Fail?

Equivalent diameter = 
$$\left(\frac{4 \times area}{\pi}\right)^{0.5}$$

Minimum Allowed Distance = 4 × Equivalent Diameter (NDO or Exhaust Point)

# **METHOD 204 ENCLOSURE DATA SHEET (cont.)**

# Near Ratio [NDO Area/Total Enclosure Area]

NDO	Surface Area (FT <sup>2</sup> )	Wall, Ceiling, or Floor Section	Surface Area (FT²)	
				NEAR ratio:
				NDOA
				NDOArea EnclosureArea =
				Pass/Fail?
TOTAL NDO AREA= TOTAL ENCLOSURE ARE			AREA=	

# **Velocity of Air through NDO (Volumetric Flow Method)**

Exhausted Air			Make	Up Air		
Exhaust Point	SCFM	Controlled? (Y/N?)	Make up point	SCFM		
					total NDO area ft <sup>2</sup>	
					Exhaust scfm - 1 make up scfm = NDO area (ft <sup>2</sup> )	—fpm
					fpm should be ≥ 200 pass/fail?	
TOTAL			TOTAL	I		

Or, Velocity of Air through NDO (Pressure Differential	Method)
Pressure Differential across the enclosure =Pressure difference > 0.007 inches H <sub>2</sub> O pass/fail?	_inches H <sub>2</sub> 0

# **METHOD 204 ENCLOSURE DATA SHEET (cont.)**

# **Direction of Air through NDO**

# **Checked by Smoke Tubes**

	Nor	mally	Direc	tion of Air Flo	w		Direction of Air Flow (One Hour Verification)						
						NDO	Initial:	10 min:	20 min:	30 min:	40 min:	50 min:	60 min:
NDO No.	Open	Closed	Into Enclosure	Out of Enclosure	Swirled	Required to be Normally Closed?	Actual Time:	Actual Time:	Actual Time:	Actual Time:	Actual Time:	Actual Time:	Actual Time:

<sup>\*</sup>Check to verify that airflow was checked at top, bottom, middle, and both sides of enclosure.

Status of doors and Are all access doo operation.		s whose	areas	are not	included	as	NDOs	closed	during	norma
Capture of VOC Em		rol (for P	TE) or t	o a point	t where it o	can	be mea	sured (f	or TTE)	
□ Yes □ No	, , , , , , , , , , , , , , , , , , ,	(	-, -, -	p				(1	,	

# **Volumetric Air Flow Calculations (Wet Bulb/ Dry Bulb)**

$$Bws = \left[\frac{e' - AP(t - t')}{P}\right]$$

e = saturated vapor presssure of water, in.Hg,

at the wet bulb temperature, t

$$A = 3.67 \times 10^{-4} \left[ 1 + 0.00064 (t' - 32) \right]$$

P = pressure, inches mercury, in the duct

t = dry bulb temperature, °F

t = wet bulb temperature, °F

Bws = water vapor in gas stream proportion

by volume

$$Md = (0.44 \times \%CO_2) + (0.32 \times \%O_2) + [0.28 \times (100 - \%CO_2 - \%O_2)]$$

$$MS = Md \times (1 - Bws) + (18 \times Bws)$$

$$Vs = \sqrt{\frac{(Ts + 460)}{Ms \times Ps}} \times \sqrt{DP} \times Cp \times 85.49$$

Acfm =  $Vs \times Area$  (of stack or duct)  $\times$  60

Dscfm = Acfm × 17.647 × 
$$\left[\frac{Ps}{(460 + Ts)}\right]$$
 × (1 - Bws)

$$Scfm = Acfm \times 17.647 \times \left[ \frac{Ps}{(460 + Ts)} \right]$$

$$Scfh = Scfm \times 60 \frac{min}{hr}$$

acfm = actual cubic feet per minute dscfm = dry standard cubic feet per minute scfm = standard cubic feet per minute scfh = standard cubic feet per hour Cp = pitot tube correction factor Ps = absolute flue gas pressure

Ms = molecular weight of gas (lb/lb mole)

Md = dry molecular weight of gas

(lb/lb mole)

### MOISTURE CALCULATIONS

$$\begin{split} V_{wc(std)} &= \frac{(V_f - V_i) \, \rho_w \, R \, T_{std}}{P_{std} M_w} = 0.04707 (V_f - V_i) \\ V_{wsg(std)} &= \frac{(W_f - W_i) \, R \, T_{std}}{P_{std} \, M_w} = 0.04715 \, (W_f - W_i) \\ V_{m(std)} &= 17.64 \, V_m \, Y \frac{P_{bar} + \frac{\Delta H}{13.6}}{T_m} \\ B_{ws} &= \frac{V_{wc(std)} + V_{wsg(std)}}{V_{wc(std)} + V_{wsg(std)}} \end{split}$$

#### Where:

B<sub>ws</sub> = Water vapor in gas stream, proportion by volume

M<sub>w</sub> = Molecular weight of water, 18.015 lb/lb-mole

P<sub>bar</sub> = Barometric pressure at the testing site, in. Hg

P<sub>std</sub> = Standard absolute pressure, 29.92 in. Hg

R = Ideal gas constant, 0.048137 (in. Hg)(ft<sup>3</sup>)/(g-mole)( $^{\circ}$ R) = [21.8348(in. Hg)(ft<sup>3</sup>)/(lb-mole)( $^{\circ}$ R)]/453.592 g-mole/lb-mole

T<sub>m</sub> = Absolute average dry gas meter temperature, °R

T<sub>std</sub> = Standard absolute temperature, 528 °R

V<sub>f</sub> = Final volume of condenser water, ml

 $V_i = Initial volume of condenser water, ml$ 

 $V_m = Dry gas volume measured by dry gas meter, dcf$ 

V<sub>m(std)</sub> = Dry gas volume measured by dry gas meter, corrected to standard conditions, scf

V<sub>wc(std)</sub> = Volume of condensed water vapor, corrected to standard conditions, scf

V<sub>wsg(std)</sub> = Volume of water vapor collected in silica gel, corrected to standard conditions, scf

 $W_f$  = Final weight of silica gel, g

W<sub>i</sub> = Initial weight of silica gel, g

Y = Dry gas meter calibration factor

 $\Delta H = Average pressure exerted on dry gas meter outlet by gas sample bag, in. H<sub>2</sub>O$ 

 $\rho_w =$  Density of water, 0.9982 g/ml

13.6 = Specific gravity of mercury (Hg)

 $17.64 = T_{std}/P_{std}$ 

 $0.04707 = ft^3/ml$   $0.04715 = ft^3/g$ 

# ppmv to lb/hr Conversion Calculations

1. ppm 
$$SO_2 \times 1.660 \times 10^{-7} = \frac{lbs/SO_2}{scf}$$

$$\frac{\text{lbsSO}_2}{\text{scf}} \times \frac{\text{scf}}{\text{min}} \times \frac{\text{60min}}{\text{hr}} = \frac{\text{lbsSO}_2}{\text{hr}}$$

2. ppmNO<sub>x</sub> ×1.194×10<sup>-7</sup> = 
$$\frac{lbs/NO_x}{scf}$$

$$\frac{\text{lbsNO}_{x}}{\text{scf}} \times \frac{\text{scf}}{\text{min}} \times \frac{60\text{min}}{\text{hr}} = \frac{\text{lbsNO}_{x}}{\text{hr}}$$

3. ppm CO 
$$\times 7.266 \times 10^{-8} = \frac{lbs/CO}{scf}$$

$$\frac{\text{lbs CO}}{\text{scf}} \times \frac{\text{scf}}{\text{min}} \times \frac{60 \text{ min}}{\text{hr}} = \frac{\text{lbs CO}}{\text{hr}}$$

4. ppm 
$$C_3H_8 \times 1.142 \times 10^{-7} = \frac{lbsC_3H_8}{scf}$$

$$\frac{\text{lbsC}_3\text{H}_8}{\text{scf}} \times \frac{\text{scf}}{\text{min}} \times \frac{60\text{min}}{\text{hr}} = \frac{\text{lbsC}_3\text{H}_8}{\text{hr}}$$

5. ppm 
$$CH_4 \times 4.164 \times 10^{-8} = \frac{IbsCH_4}{scf}$$

$$\frac{\text{lbsCH}_4}{\text{scf}} \times \frac{\text{scf}}{\text{min}} \times \frac{60\text{min}}{\text{hr}} = \frac{\text{lbsCH}_4}{\text{hr}}$$

#### **Procedures for Calibration**

## **Temperature Sensing Devices**

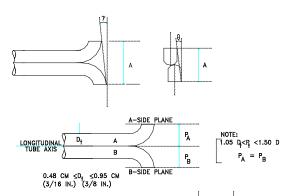
The potentiometer and thermocouples are calibrated utilizing a NBS traceable millivolt source.

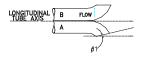
#### **Pitot Tubes**

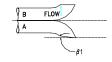
The pitot tubes utilized during this test program are manufactured according to the specification described and illustrated in the *Code of Federal Regulations*, Title 40, Part 60, Appendix A, Methods 1 and 2. The pitot tubes comply with the alignment specifications in Method 2, Section 4; and the pitot tube assemblies are in compliance with specifications in the same section.

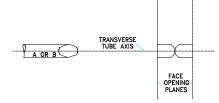
#### S TYPE PITOT TUBE INSPECTION FORM

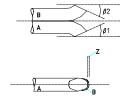
Date: 10/10/2011 Pitot Tube Nc 1 Inspectors Name: DTEM

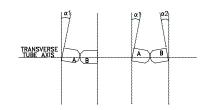


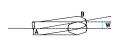












Pitot tube assembly level?

Pitot tube openings damaged? \_\_\_\_yes (explain below)

$$a_1 = 1$$
 ° (<10°),  $a_2 = 1$  ° (<10°)

$$z = A \sin g = \frac{0.008}{(in.)}; (<0.125 in.)$$

$$b_1 = 0$$
 ° (<5°),

$$b_2 = 2$$
 ° (<5°)

$$w = A \sin q = 0.025$$
 (in.); (<0.03125 in.)

$$\gamma = 0.5^{\circ}, \ \theta = 1.5^{\circ}, A = 0.938 \ \text{(in.)}$$
  $P_A = 0.477 \ \text{(in.)}, P_B = 0.477 \ \text{(in.)}, D_t = 0.375 \ \text{(in.)}$ 

$$P_A = 0.477$$
 (in.).  $P_B =$ 

0.477 (in.), 
$$D_t =$$

Calibration required?

\_\_\_\_\_yes \_\_\_x\_\_no

Dry Gas Meter No.	CM1	Date:	December 28, 2011
Standard Meter No.	16745468	Calibrated By:	BWH
Standard Meter (Y)	1.0000	Barometric Pressure:	29.65

		0 :5			la. , ,,,,					1		
		Orifice		1 1		,		Dry Gas Meter	<b>-</b>	<b>-</b> .		
		Setting in H <sub>2</sub> O	Gas Volume	Gas Volume	Temp. F°			Avg. Temp. F°	Time	Time		
Run Numb	er	Chg (H)	vr	vd	tr	tdi	tdo	td	Min	Sec	Y	Chg (H)
			П	Т	Г	1		1				
Final			35.289	61.905	59	62	62					
Initial		•	29.196	55.737	59	62	62					
Difference	1	0.20	6.093	6.168	59	62	62	62	21	30	0.993	1.373
Final			41.434	68.107	59	63	62					
Initial			35.447	62.063	59	62	61					
Difference	2	0.50	5.987	6.044	59	63	62	62	13	0	0.995	1.300
Final			47.503	74.269	58	64	62					
Initial			41.698	68.395	59	63	62					
Difference	3	0.70	5.805	5.874	59	64	62	63	10	50	0.995	1.340
Final			53.142	79.947	58	65	62					
Initial			47.691	74.437	59	64	62					
Difference	4	0.90	5.451	5.510	59	65	62	63	9	0	0.996	1.347
Final			58.938	85.809	58	63	62					
Initial			53.202	80.108	59	64	62					
Difference	5	1.20	5.736	5.701	59	64	62	63	7	45	1.011	1.204
Final		•	29.095	55.645	59	63	62					
Initial			23.404	49.895	59	62	62					
Difference	6	2.00	5.691	5.750	59	63	62	62	6	30	0.991	1.438

Average 0.997 1.333

#### Stack Temperature Sensor Calibration

Meter Box # : CM1 Name : BWH

Calibrator Model #: CL23A

Serial #: T-276953

Date Of Certification: January 11, 2011

Primary Standards Directly Traceable National Institute of Standards and Technology (NIST)

Reference	Test			
Source	Thermometer	Temperature		
Temperature (° F)	Temperature (° F)	Difference %		
0	0	0.0		
250	249	0.0		
600	599	0.0		
1200	1201	0.0		

# **Method 25A Calibration Summary**

Project:	Date:	
Client:	Operator:	
Location:	Analyzer ID:	
	Analyzer Range:	

Cal Run	Cal Level	Test Location	Cylinder ID Serial Number	Cal Gas Type	Cal Time	Expected Cal Value	Actual Response	Difference (% of cal value)	Drift (% of span)	Cylinder Pressure
	Zero							,	N/A	
D 4	Low								N/A	
Pre 1	Mid								N/A	
	High								N/A	
	Zero									
Post 1/ Pre	Low									
2	Mid									
_	High									
	Zero									
Post	Low									
2/ Pre 3	Mid									
	High									
	Zero									
Post 3	Low									
Post 3	Mid									
	High									
	Zero									
	Low									
	Mid									
	High									
	Zero									
	Low									
	Mid									
	High									

# **CALIBRATION SUMMARY**

Project Number:	Date:
Client:	Operator:
Test Location:	Box Truck:

Analyzer Type, S/N, and Span	Cal Level	Cylinder ID Serial Number	Expected Cal Value	Actual Response	Difference As % of Span	Cylinder Pressure (psi)	Cylinder Expiration Date
NO <sub>x</sub>	Zero						
	Mid						
	High						
CO <sub>2</sub>	Zero						
	Mid						
	High						
O <sub>2</sub>	Zero						
	Mid						
	High						